Effect of Predatory Stress Before Fertilization on Body Weight and Pentylentetrazole-Induced Seizure in Rat Offspring

Ehsan Saboory1, Maryam Mahmoodkhani2, Shiva Roshan-Milani2, Yousef Rasmii2

ABSTRACT

Background & Objective: Spermatogenesis and oogenesis is a primary epigenetic element that can cause lifelong modifications in neural system functions. This study was designed to elucidate whether preconceptional stress during spermatogenesis and oogenesis affects the seizure behaviors and body weight of the offspring later in life.

Materials & Methods: In this experimental study, 16 male and 32 female adult rats were included. Half of the rats were subjected to predatory stress. Then, they were coupled as follows: both subjects were intact as the control group (MC-FC), both subjects were stressed (MS-BS), male control/female stressed (MC-FS), and male stressed/female control (MS-FC). The pups were weighed and assessed for pentylentetrazole-induced seizures. In addition, aniline blue staining was performed to study the chromatin maturity of the sperm in male parents immediately after copulation.

Results: The latency of the first seizure behavior significantly diminished in stressed pups, and the duration of focal and generalized seizures significantly increased in the stressed rats. In addition, the body weight of the pups decreased in the stress groups compared to control rats, and a significant decrease was detected in the chromatin maturity of sperms in stressed rats.

Conclusion: Stress during spermatogenesis and oogenesis can be critical for the general growth (body weight) and seizure susceptibility of the offspring. Therefore, to improve reproductive outcomes, stress-lowering interventions are better to be started before conception.

Keywords: Body weight, Preconceptional, Stress disorder, Rats, Seizures

Introduction

Organisms naturally face perturbations threatening their immediate survival possibly affecting their main life-history features (1). The quality of the social environment may have considerable effects on the structure and function of nervous systems with implications for several behavioral and physiological responses (2, 3). One of the most important environmental factors is stress which is unavoidable and experienced by all and can be psychological, physical, or both (4). Stressors happening through the critical stages of development, for instance, the perinatal period, may disturb physiological functions such as metabolism, reproduction system, and growth (5). In humans, the influence of early-life stress on the brain structure and function has been revealed to last a long time after stress exposure, possibly up to adulthood. Exposure to early-life stress raises the risk of neuropsychiatric illnesses, including epilepsy (6). Stress not only increases seizure susceptibility at times of stress but also the risk of the development of epilepsy, especially in persons who have experienced early-life stress. It has been reported that prenatal stresses (PS) potentiate seizure behaviors and raise the vulnerability to seizures in rat offspring (7). In addition to the effects of PS on the first generation, several studies have mentioned the transfer of stress to the second generation (6). One likely explanation for the persistence of dynamic changes in these systems in the response to environmental factors is the participation of epigenetic mechanisms (2). Epigenetics has been used more broadly to refer to any change in gene function not associated with sequence variation. Epigenetic processes essentially influence brain development. Insults in maturation throughout pregnancy might be expected to more broadly re-program the epigenome; if they are incorporated into the germ cells, their effects will possibly become trans-generational (2). Furthermore, epigenetic characters generated within germ cells due to environmental influences all over life can shape future generations long before conception occurs (2). The connection between epigenetics and human
reproduction characterizes a very interesting field of study, generally due to the possible trans-generational effects related to epigenetic modifications of male and female gametes (8). The integrity of the sperm genome is vital for the accomplishment of a normal conception in addition to the typical growth of the fetus and children (9). In addition, it is proven that the quality of the oocyte determines the potential development of the embryo (6).

The present study was designed to elucidate whether pre-gestational stress during spermatogenesis and oogenesis (particularly chromatin changes in sperms) affects the seizure behaviors and body weight of the offspring later in life.

**Materials and Methods**

**Animals**

Wistar rats of both sexes (190-240 g) were purchased from the Urmia University of Medical Sciences, West Azerbaijan, Iran. The subjects were housed at a 12h light/dark cycle (light on at 7 a.m.) and the temperature of 20-24°C, and food and water were available ad libitum (6). After seven days of adaptation, the rats (16 adult males and 32 adult females) were randomly assigned into two groups (control and stressed male, n = 8 each; control and stressed female, n = 16 each) to form a combination of control and stressed groups for each sex.

**Predatory Stress Procedure**

A healthy adult cat was used for inducing predatory stress in the rats. As described in our previous study, the procedure was as follows: A metal cage (72x72x63 cm), with a cat inside, was placed in front of the rats' cages (Figure 1). The cage of the cat consisted of a metal floor with a removable hinged door and several small air holes on the side, so that the cat could be easily observed. To induce predatory stress, the rats were positioned in metal mesh cages (20x22x22 cm) with numerous holes in the lateral walls. The cages banned any direct contact between the predator (cat) and the rats but exposed the rats to other stimuli, e.g. the vision, smell, and noises made by the cat (6). The subjects were stressed at 8-9 a.m. and 4-5 p.m. (twice daily) for 50 or 15 consecutive days (males and females, respectively). Immediately after each session, the rats were moved to their home cage. The rats in the control group were transported to another room without any cat odors and handled similarly to the stressed rats but without being stressed. The stress procedure lasted for 50 days (in males) because it is the time required for a complete spermatogenesis cycle in male rats, while 15 days constituted three cycles of the estrous cycle in female rats (10).

Immediately after the stress procedure, the rats were assigned for copulation. The copulation method was similar to that explained in our previous study: Two female rats were coupled with one male rat for three days (the Trichus method) per cage as follows: Mc-Fc (both rats were control), Ms-Fc (the male was exposed to stress, but the female was control), Mc-Fs (the male rat was control but the female was stressed), and Ms-Fs (the two rats were stressed) (10).

The pregnant subjects were transferred to new cages and kept in normal conditions for the entire pregnancy period. The duration of pregnancy was the same, 21 days, in four groups. Upon delivery, the pups were counted, weighed, and gendered. All the litter was culled to eight per dam. The rat pups were allocated according to their own parents’ grouping as follows: Both fathers and mothers were control (Mc-Fc); just the mother was stressed (Mc-Fs); just the father was stressed (Ms-Fc); and both the father and mother were stressed (Ms-Fs) (10). There were 64 pups in each group (8 dams x 8 pups = 64).

**Sperm Sampling in Adult Rats**

Under anesthesia with isoflurane, the stressed and control male parents were subjected to sperm sampling in the morning (n = 8 per group) after copulation. Both epididymides of each rat (at the caudal region) were transferred to a glass Petri dish containing an 80% DMEM/F12 (DMEM/F12; ATCDHF, St. Louis, USA) culture supplemented with a 20% fetal bovine serum (FBS; Sigma, St. Louis, USA) medium pre-warmed to 37°C. The epididymides were crushed, and six cuts were made with an insulin syringe. The samples were incubated for 30 min at 37°C in 5% CO₂; then, spermatozoa were released from the epididymides (10),...
Chromatin Quality and Acidic Aniline Blue (AB) Staining

For each neat semen sample, the sperm nuclei chromatin condensation was evaluated using AB staining. Aniline Blue selectively dyes histones with abundant lysine and can demonstrate the sperm chromatin irregularities associated with histone anomalies. Fixations of air-dried smears of rat semen samples were prepared in buffered glutaraldehyde (3%) in a phosphate buffer (0.2 M and pH 7.2) for half an hour at 22-24°C. Then, the smears were stained with aqueous AB stain (5%) in acetic acid (4% and pH = 3.5) for 7 min (11). In a stained smear, 200 sperm were studied under a light microscope with x100 magnification. In this staining, the spermatooza with light-blue nuclei are considered as normal, and those with dark-blue nuclei are counted as abnormal (12).

Body Weight Measurement

Upon delivery, the rat pups for each dam were counted and weighed in the morning at 08:30 on the first postnatal day (P1). Weighing of the pups was repeated at 08:30 on postnatal days 6, 15, 25, and 30 (P6, P15, P25, and P30, respectively).

Behavioral Assessment

In the pentylenetetrazole (PTZ) model, seizures are commonly induced by a single systemic administration, and PTZ can even induce status epilepticus if given in a sufficient amount (13). On P30, one male and one female pup from each dam were subjected to seizure induced by PTZ. Then, the pups were intraperitoneally (IP) injected with PTZ (55 mg/kg) (6, 14). After injection, the behavior of each rat was assessed for one hour with a digital camera and direct observation. The behavioral rating was evaluated by means of a previously defined scale: normal behaviors = 0; immobilization with or without sniffing = 1; forelimb clonus with head nodding named as short myoclonic jerk = 2; tail extension and nonstop myoclonic jerk = 3; widespread limbic seizures with kangaroo position = 4; and nonstop tonic or tonic-clonic seizures (15, 16). Other variables were also evaluated as follows:

- Time to the onset of the first epileptic behavior (sec): time interval between the injection of PTZ and the appearance of the first seizure; the number of focal seizures: the number of focal seizures in 60 minutes (e.g. leg opening, immobility, facial twitching, …); duration of focal seizure (min), total time of all focal seizures in 60 minutes; the number of tonic-clonic seizures, the number of tonic-clonic seizures in 60 minutes that each rat experienced; duration of tonic-clonic seizure (sec), total time of all tonic-clonic seizures in 60 minutes

Ethical Consideration

This experimental study was approved by the Ethics Committee in West Azerbaijan Province, Iran (ethical code: IR.1395.229).

Statistical Analysis

Data distribution was checked by the Kolmogorov-Smirnov test. The normally distributed data were analyzed using one-way ANOVA (body weight and epileptic behaviors, except for the number of focal seizures and the onset of the first seizure) followed by Tukey's post-hoc test when required. The data related to the latency (onset) of the first seizures and the number of focal seizures were evaluated by Kruskal-Wallis or Mann-Whitney U test. Also, t-test was performed for the comparison of chromatin maturity between stressed and control rats. The findings were stated as mean ± standard error of the mean (SEM), and the P-value < 0.05 was considered significant.

Results

Effects of Pregestational Stress on Seizure

On P30, PTZ-induced seizure was assessed in rat pups. First, the data of the pups for male and female pups were independently analyzed; no significant difference was detected between the two sexes. Then, the data related to the seizure of male and female pups were combined and evaluated together (Table 1).

Effects of Pre-conception Stress on Bodyweight in Rat Pups

There were significant differences in body weight among the groups at different time points (Table 2). In pregestational stress, the mean body weight of the offspring significantly decreased at birth (P1, P=0.03) and P6 (P=0.02) compared to the control pups.

Effects of Stress During Spermatogenesis on Chromatin Maturity

Remarkable differences were found in the chromatin maturity of sperms between stressed and control adult rats (Figure 2). Also, an illustration of sperms with normal and abnormal chromatin maturity is shown in Figure 3.

![Figure 2. Effect of stress during spermatogenesis on the chromatin maturity of sperms in adult rats; *indicates P=0.004 with the stress group.](image-url)
Effect of Predatory Stress Before Fertilization

Figure 3. Aniline blue staining of sperms in adult Wistar rat subjected to predatory stress for 50 consecutive days; (A) a normal sperm; (B and C) sperms with abnormal maturity (×1000).

Table 1. Effect of pre-conception stress during spermatogenesis and oogenesis on PTZ-induced seizure in the rat pups

<table>
<thead>
<tr>
<th>Epileptic behaviors</th>
<th>Pups grouping</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>MC-FC</td>
<td>MC-FS</td>
</tr>
<tr>
<td>Time to onset of first epileptic behavior</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(Sec)</td>
<td>119.87±14.61</td>
<td>#18.25±1.95</td>
</tr>
<tr>
<td></td>
<td>#P&lt;0.01 vs. Mc-Fc</td>
<td></td>
</tr>
<tr>
<td>Number of focal seizure</td>
<td>1.5±0.32</td>
<td>2.12±0.35</td>
</tr>
<tr>
<td></td>
<td>non-significant</td>
<td></td>
</tr>
<tr>
<td>Duration of focal seizure (min)</td>
<td>9.25±2.54</td>
<td>14.62±2.01</td>
</tr>
<tr>
<td></td>
<td>*P&lt;0.04 vs. Mc-Fc</td>
<td></td>
</tr>
<tr>
<td>Number of tonic-clonic seizure</td>
<td>0.5±0.18</td>
<td>1.12±0.26</td>
</tr>
<tr>
<td></td>
<td>non-significant</td>
<td></td>
</tr>
<tr>
<td>Duration of tonic-clonic seizure (Sec)</td>
<td>1.37±0.53</td>
<td>*13.75±3.09</td>
</tr>
<tr>
<td></td>
<td>*P&lt; 0.03 vs. Mc-Fc</td>
<td></td>
</tr>
</tbody>
</table>

Data presented as mean± SE: # non-parametric Kruskal Wallis test, * Tukey post-hoc test.
MC-FC: both parents’ control, MC-FS: male control/female stressed, MS-FC: male stressed/female control, MS-FS: both parents stressed

Table 2. Effects of pre-gestational stress on bodyweight in rat pups

<table>
<thead>
<tr>
<th>Groups</th>
<th>P1</th>
<th>P6</th>
<th>P15</th>
<th>P25</th>
<th>P30</th>
</tr>
</thead>
<tbody>
<tr>
<td>MC-Fc</td>
<td>6.45 ± 0.65</td>
<td>11.59±0.22</td>
<td>23.61±0.69</td>
<td>42.49± 1.1</td>
<td>56.27± 0.95</td>
</tr>
<tr>
<td>MC-Fs</td>
<td>*6.15±.09</td>
<td>*10.36±0.37</td>
<td>21.15±0.58</td>
<td>39.93±1.43</td>
<td>53.71±0.81</td>
</tr>
<tr>
<td>Ms-Fc</td>
<td>6.38 ±0.09</td>
<td>11.24±0.12</td>
<td>21.43±0.26</td>
<td>41.43±0.47</td>
<td>53.36±0.7</td>
</tr>
<tr>
<td>Ms-Fs</td>
<td>5.45±0.11</td>
<td>*10.32±0.19</td>
<td>20.99±0.58</td>
<td>39.84±0.66</td>
<td>52.85±1.09</td>
</tr>
<tr>
<td>P-value</td>
<td>*P&lt;0.03 vs. Mc-Fc</td>
<td>*P&lt;0.001 vs. all groups</td>
<td>*P&lt;0.02 vs. Mc-Fc</td>
<td>non-significant</td>
<td>non-significant</td>
</tr>
</tbody>
</table>

Data presented as mean± SE and analyzed by One-way ANOVA and Tukey tests.
P1, postnatal day 1, P6, postnatal day 6, P15, postnatal day 15, P25, postnatal day 25, P30, postnatal day 30
MC-FC: both parents control, MC-FS: male control/female stressed, MS-FC: male stressed/female control, MS-FS: both parents stressed

Discussion

The effects of pre-gestational stress on PTZ-induced seizure, the body weight of the offspring, and the sperm chromatin changes of male parents were investigated.

The main findings were that pregestational stress caused a significant increase in generalized seizures and decreased the body weight of the offspring at birth.
(P1) and on P6. In addition, there was a significant decrease in the percent of spermatozoa with mature chromatin in the stressed rats compared to the control group.

**Effects of Pre-conception Stress on Epileptic Behaviors in Rats**

The results demonstrated the negative effects of pre-pregnancy stress on the offspring. In this regard considerable evidence exists supporting the fact that many factors can persuade hostile effects in the descendants following parental exposure and, consequently, represent a threat in this respect (17). PS can be regarded as a primary epigenetic element able to cause lifelong modifications in neural system functions (18). In previous studies, the effect of early stress on growth has been studied; early-life stress, such as prenatal stress, is reported to be proconvulsant. In one study, PS amplified seizure and diminished the latency of seizure from early infancy up to later life in a kindling epilepsy model (19). Moreover, a vital element influencing adult nervous systems (such as the hippocampus, amygdala, and the hypothalamo-pituitary-adrenal (HPA) axis) is the exposure of parents to noticeable stimuli in the environment before pregnancy (20). Notably, the trans-generational influence of the chronic stress of parents is not limited to the perinatal window; alterations in the descendants' stress-associated behavior have been stated following the exposure of parents to stress up to adulthood (21). The findings of our study are consistent with the results of previous investigations. The mechanisms by which long-lasting parental stress can disturb the descendants' neurodevelopment are not fully understood yet. A mechanism explaining the impact of pre-pregnancy stress on the offspring is the HPA axis. Environmental factors, mainly a positive stress history, contribute to the developmental programming of the HPA axis (21). Increasingly, the HPA axis is being considered as a compound physiological system able to facilitate huge variations in the physiological responses of subjects to stressors (1). It has been reported that a repetitive experience of PS-induced disruption in the function of the HPA axis and modification of its feedback regulation originate a greater basal secretion of glucocorticoid hormones (22). Furthermore, it has been indicated that the serum corticosterone and adrenocorticotropic hormone (CRH) levels were higher in maternal rats with pre-gestational chronic unpredictable stress (CUS) than in the control group, both after CUS and delivery. Chronic stress not only leads to the imbalance of the neuroendocrine network of maternal rats but also impacts the HPA axis of the fetus via the maternal-placental-fetal interface (23).

Previous studies have argued that stress before gestation will possibly affect the hippocampal maturation and impair the memory function of the offspring, and cause reduced brain-derived neurotrophic factor levels (24). Perhaps the same argument applies to the effect of pre-pregnancy stress on the offspring's epileptic behaviors. Meanwhile, it has been shown that stress disturbs the integrity of the sperm chromatin (4) which is important as it indicates that sperms possibly pass damaged genes onto the zygote and embryo which may eventually influence the health of the descendants (25). In the present study, in groups where the father was stressed, the chromatin maturation rate decreased and the health of the children was affected. Also, research on some animal species indicates that egg quality is effective on the growth and health of the children (26). Here, the stress state in the pre-pregnancy period in parents showed that stress during gametogenesis can transfer to the offspring, thereby altering the offspring's vulnerability to seizure. In conclusion, the present study demonstrated that the exposure of rats to stress during spermatogenesis and oogenesis can change their next generation's vulnerability to PTZ-induced epileptic behaviors later in life.

**Effects of Pre-conception Stress on the Body Weight of the Offspring**

Studies have reported that PS leads to a low birth weight. Animal studies have demonstrated that exposing animals (such as rats and mice) to stressors during pregnancy is associated with lower birth or fetal weights. Maternal stress affects the growth and organ development of fetuses, targeting some organs more than others, strikingly attenuating glucose transporter expression, and diminishing fetal plasma glucose, growth hormone, and CRH levels (27). There have been a few experiments examining the effect of stress before pregnancy on the pups’ body weight. According to the findings, a concomitant decrease in the weight of dams and their fetuses indicates that growth retardation will possibly relate to the dams with CUS before pregnancy (24). As our study showed, in the case of mothers exposed to stress during oogenesis, their offspring showed a lower body weight on P1 and P6. Ronnenberg et al. studied the connection between pre-gestation body mass index and delivery outcomes among women in China; children born to women who were underweight before gestation were at a higher risk for fetal growth retardation (28). Another study assessed the welfare of prescribing iodized oil to women before pregnancy or during early pregnancy in Algeria. The children of treated mothers had a significantly greater birth weight compared to control women (29). These studies, in line with our study, show that maternal conditions before pregnancy can affect the health and weight of the offspring. On the other hand, the findings of another study identified no difference in the body weight of the offspring of subjects undergoing pre-gestational stress and the controls (24), a finding that contrasts those of the present study. Also, it has been reported that sexual behavior can be the most susceptible feature of male reproduction to stress due to the antagonistic association between male sex hormone and glucocorticoids. This principle of sperm quality might predict the treatment success as recommended by its links at several issues in the reproductive course, such as interrupted
preimplantation of the embryo and birth defects in the children (30). The mechanisms by which pregestational stress affects pup growth remain largely unknown. Nevertheless, pregestational CUS leads to a remarkable rise in GC such as corticosterone levels in the offspring of the stressed subjects (24). Also, an animal study suggests that periconceptional undernutrition may influence the HPA axis which, in turn, influences the outcomes (28). However, the long-term influence of stress has been connected to a disruption in the HPA axis physiology and a change in its negative feedback regulation, affecting fetal growth and development (22). Our findings indicated that in the case of mothers exposed to stress during oogenesis, their offspring showed a lower body weight at birth and six days after birth. This result is consistent with the findings of the above-mentioned studies, suggesting that parental stresses before conception affect the birth weight and early neonatal body weight of the newborns (10). Meanwhile, to the best of our knowledge, this study is unique because both parents’ stress and the single-parent (maternal or paternal) stress before conception were investigated on seizure susceptibility and body weight in the offspring, whereas previous studies have been conducted on single parents (mother or father alone).

Effects of Pre-conception Stress on Sperm Chromatin Integrity

The successful fertilization of eggs and subsequent development of the offspring will greatly depend on the quality of gametes produced by the parents. A previous study has indicated that stress has a negative impact on various parameters (such as sperm morphology and motility) associated with semen quality (10). A growing body of evidence connects the DNA damage of the sperm with mutation-related hazards and disorders in the offspring (31). Additionally, researchers believe that elevated psychological stress can be linked with amplified oxidant agents’ synthesis, and chronic stress exposure can raise the production of reactive oxygen species (ROS) (32). It has been shown that oxidative stress affects sperm chromatin integrity and leads to single- and/or double-strand DNA breaks (33). Meanwhile, sperms are not able to reinstate the oxidative stress-induced impairment since they lack the essential enzymatic repair systems. This weakness is a criterion which makes the spermatozoa very vulnerable to oxidative damage (33). Defects in the male genome have been shown to lead to post-fertilization failure (34). Previous studies recommend that DNA-fragmented sperm can cause several reproduction-related problems such as conception failure, abortions, and malformations (32). Thus, the findings of the present study confirmed the result of previous studies showing that pregestational stresses disturb the maturity of the chromatin of sperms in stressed subjects (32,33). In conclusion, a precise assessment of the stress status and seminal ROS levels should be considered as an integral part of the reproduction work-up of men in order to assist clinicians in clarifying the fertility status and thus providing the best treatment regime for patients.

Conclusion

Parents’ stress during spermatogenesis and oogenesis can be critical for the health of the offspring. One of the probable reasons for these findings is the sensitivity of the gametes, in particular, male gametes, to environmental factors. The findings suggest that alterations caused by environmental factors in the gametes can affect the health of the children as epigenetics effects. We tried to show that the stress of spermatogenesis can affect the chromatin maturation of the sperm nucleus. It reduces the quality of the sperm and promotes the transfer of defective genes to the offspring. Therefore, to improve reproductive outcomes, stress-lowering interventions are better to be started before conception.

Acknowledgments

This study was supported by the Neurophysiology Research Center, Urmia University of Medical Sciences, Urmia, Iran.

Conflict of Interest

Authors declared no conflict of interest.

References


Effect of Predatory Stress Before Fertilization


How to Cite This Article:

Download citation: BibTeX | RIS | EndNote | Medlars | ProCite | Reference Manager | RefWorks

Send citation to: Mendeley | Zotero | RefWorks | RefWorks